

Specifically, the Examiner states that the term "method of claim 0" renders the claim indefinite.

In response thereto, Claim 8 is hereby amended to overcome this rejection, as suggested by the Examiner, to recite dependence upon Claim 7.

In addition, the Examiner states that Claim 8 is also indefinite when  $R^2$  is H.

In response thereto, Claim 8 is hereby amended to overcome this rejection by defining the term  $R^2$ , as used therein, as not including "H" as one of ordinary skill in the art would clearly understand that a compound of formula III does not require treatment with  $R^2$ -L, to form a compound of formula II, wherein  $R^2$  is H, as, in this particular instance, this formula III compound is the same compound as the formula II.

Thus, in view of the amendment, the rejection of Claim 8 is now moot.

#### Conclusion

Applicants concur with the Examiner's decision that Claims 1-7 and 9-10 are allowable.

Regarding Claim 8, based on the foregoing, Applicant respectfully submits that the Examiner's rejection, under 35 USC 112, second paragraph, is now moot. Therefore,

Applicant respectfully requests that the rejection of  
Claim 8 under 35 USC 112, second paragraph, be withdrawn.  
Applicant further requests that a notice of allowance be  
issued for pending Claims 1-10.

Respectfully Submitted:

Date: 14 April 2003



Scott Alexander McNeil  
Attorney for Applicant  
Registration No. 37,185

Pfizer Inc.  
Patent Department, Box 519  
Eastern Point Road  
Groton, Connecticut 06340  
(860) 715-0871

## ATTACHMENT TO AMENDMENT

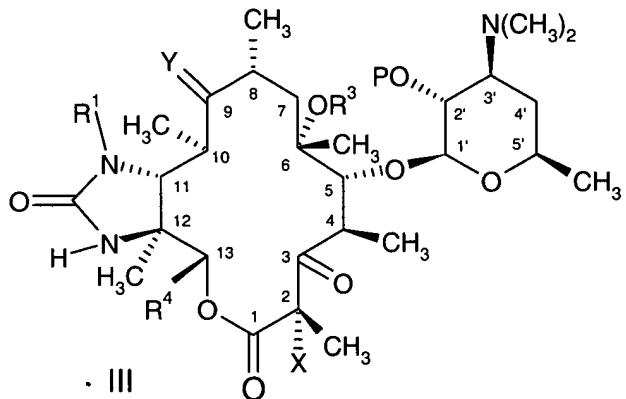
Version with Markings to Show Changes Made

DO NOT ENTER THIS VERSION

CLAIMS

2(twice amended). The compound of claim [0] 1 wherein Y is =O or =NOR<sup>5</sup>, R<sup>1</sup> is (4- to 10-membered heterocyclic) C<sub>1</sub>-C<sub>6</sub> alkyl, wherein the heterocyclic is substituted by 4- to 10-membered heterocyclic, R<sup>2</sup> is C<sub>1</sub>-C<sub>10</sub> alkyl or C<sub>2</sub>-C<sub>10</sub> alkenyl, R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, R<sup>4</sup> is ethyl, R<sup>5</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, and R<sup>6</sup> is H.

8(twice amended). The method of claim [0] 7 further wherein the compound of formula II is prepared by treating a compound of the formula



with a strong base and a compound of formula R<sup>2</sup>-L, where L is a leaving group, and wherein R<sup>2</sup> is selected from the group consisting of [H,] C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, (4- to 10-membered heterocyclic) C<sub>1</sub>-C<sub>6</sub> alkyl, (4- to 10-membered

heterocyclic) C<sub>2</sub>-C<sub>6</sub> alkenyl, (4- to 10-membered heterocyclic) C<sub>2</sub>-C<sub>6</sub> alkynyl, (C<sub>6</sub>-C<sub>10</sub> aryl) C<sub>1</sub>-C<sub>6</sub> alkyl, (C<sub>6</sub>-C<sub>10</sub> aryl) C<sub>2</sub>-C<sub>6</sub> alkenyl, and (C<sub>6</sub>-C<sub>10</sub> aryl) C<sub>2</sub>-C<sub>6</sub> alkynyl wherein said alkyl moieties of the foregoing groups are optionally substituted by halo or C<sub>1</sub>-C<sub>6</sub> alkyl, and wherein said heterocyclic moieties are optionally substituted by 4- to 10-membered heterocyclic, (4- to 10-membered heterocyclic) C<sub>1</sub>-C<sub>6</sub> alkyl, or (C<sub>6</sub>-C<sub>10</sub> aryl) C<sub>1</sub>-C<sub>6</sub> alkyl, and further wherein the aryl and heterocyclic moieties of each of the foregoing groups and optional substituents is optionally substituted by 1 to 4 R<sup>7</sup> groups.